

In the Claims:

Amend claims 23, 25, 27, 29, 30, 31, 33, 41 and 42 as follows:

Sub D1

23. (Amended) A method for detecting prostate cancer in a patient comprising:
(a) obtaining a biological sample from the patient;
(b) contacting the sample with at least two oligonucleotide primers in a polymerase chain reaction, wherein at least one of the oligonucleotides is specific for a DNA molecule comprising a sequence selected from the group consisting of SEQ ID NO:110 and complements of SEQ ID NO:110; and
(c) detecting in the sample a DNA sequence that amplifies in the presence of the oligonucleotide primers thereby detecting prostate cancer, wherein the biological sample is selected from the group consisting of: blood and [serum] semen.

Sub D2

25. (Amended) A method for detecting prostate cancer in a patient comprising:
(a) obtaining a biological sample from the patient;
(b) contacting the sample with at least two oligonucleotide primers in a polymerase chain reaction, wherein at least one of the oligonucleotides is specific for a DNA molecule comprising a sequence selected from the group consisting of SEQ ID NO:111 and complements of SEQ ID NO:111; and
(c) detecting in the sample a DNA sequence that amplifies in the presence of the oligonucleotide primers thereby detecting prostate cancer, wherein the biological sample is selected from the group consisting of: blood and [serum] semen.

Sub D5

27. (Amended) A method for detecting prostate cancer in a patient comprising:
(a) obtaining a biological sample from the patient;
(b) contacting the sample with at least two oligonucleotide primers in a

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C3
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polymerase chain reaction, wherein at least one of the oligonucleotides is specific for a DNA molecule comprising a sequence selected from the group consisting of SEQ ID NO:115 and complements of SEQ ID NO:115; and

(c) detecting in the sample a DNA sequence that amplifies in the presence of the oligonucleotide primers thereby detecting prostate cancer, wherein the biological sample is selected from the group consisting of: blood and [serum] semen.

29. (Amended) A method for detecting prostate cancer in a patient comprising:

(a) obtaining a biological sample from the patient;

(b) contacting the sample with at least two oligonucleotide primers in a polymerase chain reaction, wherein at least one of the oligonucleotides is specific for a DNA molecule comprising a sequence selected from the group consisting of SEQ ID NO:173-175, 177 and complements of SEQ ID NO:173-175 and 177; and

(c) detecting in the sample a DNA sequence that amplifies in the presence of the oligonucleotide primers thereby detecting prostate cancer, wherein the biological sample is selected from the group consisting of: blood and [serum] semen.

30. (Amended) The method of claim 29, wherein at least one of the oligonucleotide primers comprises at least about 10 contiguous nucleotides of a DNA molecule comprising a sequence selected from the group consisting of SEQ ID NO:173-175 and 177.

31. (Amended) A method for detecting prostate cancer in a patient comprising:

(a) obtaining a biological sample from the patient;

(b) contacting the sample with at least two oligonucleotide primers in a polymerase chain reaction, wherein at least one of the oligonucleotides is specific for a DNA molecule comprising a sequence selected from the group consisting of SEQ ID NO:223 and complements of SEQ ID NO:223; and

(c) detecting in the sample a DNA sequence that amplifies in the presence of

CL
nt the oligonucleotide primers thereby detecting prostate cancer, wherein the biological sample is selected from the group consisting of: blood and [serum] semen.

Su 33. (Amended) A method for detecting prostate cancer in a patient

Su *Da* comprising:

Su (a) obtaining a biological sample from the patient;

Su (b) contacting the sample with at least two oligonucleotide primers in a polymerase chain reaction, wherein at least one of the oligonucleotides is specific for a DNA molecule comprising a sequence selected from the group consisting of SEQ ID NO:224 and complements of SEQ ID NO:224; and

CS (c) detecting in the sample a DNA sequence that amplifies in the presence of the oligonucleotide primers thereby detecting prostate cancer, wherein the biological sample is selected from the group consisting of: blood and [serum] semen.

Su 41. (Amended) A method for detecting the presence of a DNA molecule

Su *Da* comprising [SEQ ID NO:115] a sequence selected from the group consisting of: SEQ ID NO:173-175 and 177 in a biological sample, the method comprising:

CL (a) contacting the sample with at least two oligonucleotide primers in a polymerase chain reaction, wherein at least one of the oligonucleotides is specific for a DNA molecule comprising a sequence selected from the group consisting of: SEQ ID NO:173-175 and 177; and

(b) detecting in the sample a DNA sequence that amplifies in the presence of the oligonucleotide primers.

42. (Amended) The method of claim [39] 41, wherein at least one of the

oligonucleotide primers comprises at least about 10 contiguous nucleotides of a DNA molecule comprising a sequence selected from the group consisting of: SEQ ID NO: 173-175 and 177.